

### **EMPOWERING INNOVATIONS**

## IVT RNA Raw Materials

FOR CUTTING-EDGE RESEARCH AND ADVANCEMENTS

The first successful report of the use of in vitro transcribed (IVT) mRNA in animals was published in 1990 after reporter gene mRNAs were injected into mice. Scientists have been using the in vitro transcribed (IVT) RNA approach to create therapies and vaccines for decades due to its short development time, high safety, high specificity, and simple manufacturing.

### The Application of IVT RNA Technology



**Therapeutics** 



Vaccine



**Drug Discovery** 



**Basic Research** 

- CRISPR/ Cas Gene Editing
- In Vivo Production of Secretable Protein/ Antibody
- Reprogramming Cells

### The Workflow of **IVT RNA Preparation**



Target Design and DNA **Template Generation** 



Plasmid Production, Purification, and Linearization



**RNA Synthesis** 



**RNA Purification** 



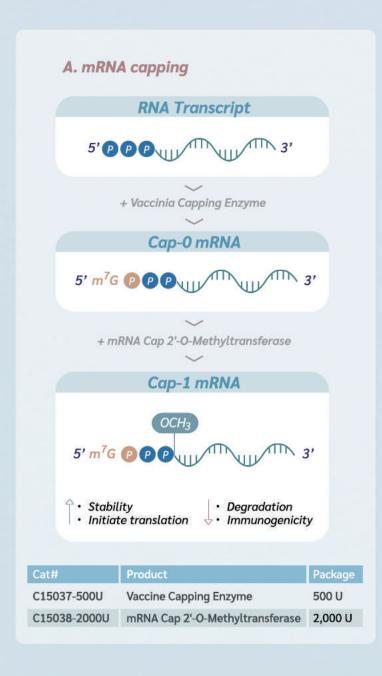
**Analytics** 



**Formulation** 



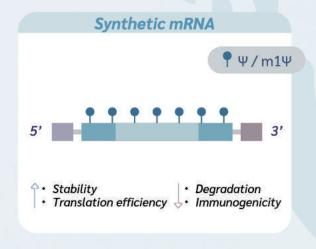
Lab Use/ **Industrial Manufacturing** 



## Key Elements for Improving the Performance of mRNA

### B. Modified nucleotide

Pseudouridine ( $\Psi$ ) and N1-Me-pUTP (m1 $\Psi$ ) can be used to replace uridine in the IVT mRNA. It is demonstrated that the modified UTP can enhance RNA stability and decrease anti-RNA immune response.



Cat#	Product	Package
C15040-100UL	Pseudo UTP Sodium Solution	100 µL
C15041-100UL	N1-Me-pUTP Sodium Solution	100 μL

## The Critical Quality Management of mRNA Purity - dsRNA Detection Assay

Cat#	Product	Package
C15039-200UG	Anti-dsRNA antibody [clone J2]	200 μg

When developing IVT mRNA-based therapies or vaccines, high-quality and purity in vitro transcribed mRNA is of the utmost importance. Double-stranded RNA (dsRNA) impurities are one of the extremely concerning byproducts because they would inhibit the synthesis of the antigen protein and trigger an unfavorable immunological response.

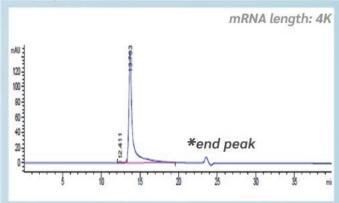
# dsRNA Amount Exposure • 2.5 ug • 1.3 ug • 0.6 ug • 0.3 ug 20s 10s 5s 1s

#### **Features**

- · High Sensitivity
- · High Specificity
- · Cost-effective

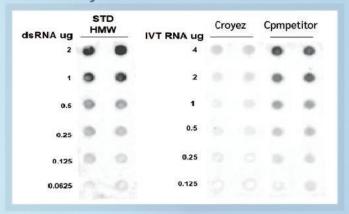
### **High Quality Final mRNA Products** by Using Croyez's Raw Materials

### mRNA purity by SEC-HPLC



Croyez's final mRNA product has a extremely high purity.

### dsRNA Analysis



Croyez's final mRNA product has a significantly lower concentration of dsRNA byproducts compared to the competitor's product.

### Comprehensive **Solutions** for IVT RNA **Production**

- · Generate high quality and high purity **IVT mRNA**
- · Flexible formulation buffers to assist you find the optimal condition

Cat#	Product	Package	
C15009-K01	NTP Set, 100 mM Solutions	1 mL*4	
C15010H-25000U	T7 RNA Polymerase (200 U/ μL)	25,000U	
C15027-K01	T7 RNA Polymerase Transcription Buffer Set	10,000 U	
C15027-K02	*Include buffer A~I (200 U/ µL)	25,000 U	
C15010HA-25000U	T7 RNA Polymerase with specific buffer A(200 U/ $\mu$ L)		
C15010HB-25000U	T7 RNA Polymerase with specific buffer B (200 U/ $\mu$ L)	00 U/ μL)	
C15010HC-25000U	T7 RNA Polymerase with specific buffer C (200 U/ $\mu$ L)		
C15010HD-25000U	T7 RNA Polymerase with specific buffer D (200 U/ $\mu$ L)		
C15010HE-25000U	T7 RNA Polymerase with specific buffer E (200 U/ $\mu$ L)	1000	
C15010HF-25000U	T7 RNA Polymerase with specific buffer F (200 U/ $\mu$ L)		
C15010HG-25000U	T7 RNA Polymerase with specific buffer G (200 U/ μL)		
C15010HH-25000U	T7 RNA Polymerase with specific buffer H (200 U/ µL)		
C15010HI-25000U	T7 RNA Polymerase with specific buffer I (200 U/ $\mu$ L)		
C15022-1ML	ATP Solution (100 mM)	1 mL	
C15023-1ML	UTP Solution (100 mM)	1 mL	
C15024-1ML	CTP Solution (100 mM)	1 mL	
C15025-1ML	GTP Solution (100 mM)	1 mL	
C15026-10U	Inorganic Pyrophosphatase (Yeast)	10U	
C15026-50U		50U	





